

K004015

Premarket Notification [510(k)] Summary
Tab 4

December 15, 2000

MAR 13 2001

Trade Name: Beam Block Trays

Common Name: Block support trays for Radiation Therapy

Classification Name: Medical Linear Accelerator Accessory, 90 IYE (per 21 CFR section 892.5050)

Manufacturer's Name: Arplay Medical S.A.
Address: 1 Route de Citeaux
21110 Izeure
France

Corresponding Official: Richard Borgi, MD
Title: President and CEO
Telephone: +33-3-8029 7401
Fax: +33-3-8029 7622

Predicate: Aktina Medical Physics Corp. Beam Block Trays, K964840

Device Description: The Beam Block Trays are accessories intended to support beam blocks used to define the radiation field during radiation therapy. They are designed and machined to meet the individual specifications of the tray mounts of all brands and models of linear accelerators and cobalt treatment machines.

The Beam Block Trays are fabricated with and without holes or slots for beam defining block attachment to meet the customer's requirements. They are also supplied with and without coding as required by the manufacturer's tray mount and the requirements of the customer.

Intended Use: Accessory to support beam blocks used to define the radiation field during radiation therapy.



Technological Characteristics: See the attached predicate comparison table.

| # | Feature | Beam Block Trays, K964840 | Arplay Medical Beam Block Trays |
|---|-----------------|--|--|
| 1 | Materials | Acrylic | Acrylic or Polycarbonate |
| 2 | Compatilability | All models of Siemens linear accelerators | All brands and models of linear accelerators and cobalt treatment machines |
| | | | |

The Arplay Medical Beam Block Trays have the same intended use and performance characteristics as the predicate device. No new issues of safety of effectiveness are introduced by this device.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

MAR 13 2001

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Richard Borgi, M.D.
President & CEO
Arplay Medical S.A.
1 Route de Citeaux
21110 Izeure
FRANCE

Re: K004015
Beam Block Trays
Dated: December 15, 2000
Received: December 27, 2000
Regulatory Class: II
21 CFR §892.5050/Procode: 90 IYE

Dear Dr. Borgi:


We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for *in vitro* diagnostic devices), please contact the Office of Compliance at (301) 594-4639. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsma/dsmamain.html>".

Sincerely yours,


Daniel G. Schultz, M.D.
Acting Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure (s)

Tab 3

Indications For Use

510(k) Number: K004015

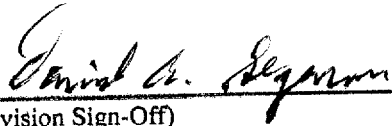
Device Name: Beam Block Trays

Indications for Use:

Accessory to support beam blocks used to define the radiation field during radiation therapy.

(PLEASE DO NOT WRITE BELOW THIS LINE)

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)

Division of Reproductive, Abdominal, ENT,
and Radiological Devices

510(k) Number K004015

Prescription Use ☒
(per 21 CFR 801.109)

OR

Over-The-Counter Use ☐